

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: April 5, 2001, 19:32:00 ; Search time 10277.7 Seconds
(without alignments)
4505.438 Million cell updates/sec

Title: US-09-513-888-1
Perfect score:
Sequence: 1 gcctccaaagacctgccc.....tgccattctcacggcctct 9048

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1118133 seqs, 2558875100 residues

Total number of hits satisfying chosen parameters: 2236266

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : GenEmbL:
1: gb_ba1: *
2: gb_ba2: *
3: gb_on: *
4: gb_ov: *
5: gb_ph: *
6: gb_pl1: *
7: gb_pl2: *
8: gb_pr1: *
9: gb_pr2: *
10: gb_pr3: *
11: gb_ro: *
12: gb_sy: *
13: gb_un: *
14: em_fun: *
15: em_hum1: *
16: em_hum2: *
17: em_in: *
18: em_om: *
19: em_or: *
20: em_ov: *
21: em_pat: *
22: em_ph: *
23: em_pl: *
24: em_ro: *
25: em_sts: *
26: em_sy: *
27: em_un: *
28: em_v1: *
29: gb_da3: *
30: gb_in1: *
31: gb_in2: *
32: gb_in3: *
33: gb_pl3: *
34: gb_pr4: *
35: em_da1: *
36: em_ba2: *
37: em_htg1: *
38: em_htg2: *
39: em_htg3: *
40: em_htg4: *
41: em_htg5: *
42: em_htg6: *
43: em_htg7: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB	ID	Description
1	9046.8		100.0	9108	10	AF123653	Homo sapi
c	2	426.8	54.5	172162	64	AC025953	Homo sapi
c	3	4199.8	46.4	5492	10	AF123559	Homo sapi
	4	807.2	8.9	1515	10	AF123556	Homo sapi
	5	807.2	8.9	1614	10	AF123555	Homo sapi
	6	807.2	8.9	1692	10	AF123557	Homo sapi
	7	807.2	8.9	1722	10	AF123558	Homo sapi
	8	346	3.8	633	10	AF123554	Homo sapi
c	9	229.2	2.5	163683	64	AC025412	Homo sapi
	10	226.2	2.5	152511	60	AC019369	Homo sapi
	11	225.6	2.5	179338	63	AC024740	Homo sapi
c	12	224.6	2.5	158440	34	AL355392	Human DNA
	13	224.2	2.5	177560	59	AC016695	Homo sapi
	14	224	2.5	227949	65	AC026803	Homo sapi
	15	223.6	2.5	160659	66	AC036335	Homo sapi
	16	223.6	2.5	180516	84	CNS010SH	
	17	223.6	2.5	190128	54	AC007374	Homo sapi
	18	223.6	2.5	213945	74	AL355337	Homo sapi
c	19	223.4	2.5	89818	8	AC002126	Homo sapi
c	20	223.2	2.5	141292	72	AF235106	Homo sapi
c	21	223	2.5	156345	62	AC023015	Homo sapi

Vassiliev, H.; Viel, R.; Vo, A.; Wilson, B.; Wu, X.; Wyman, D.; Ye, W.-J.,
 Young, G.; Zainoun, J.; Zimmer, A. and Zody, M.
 Direct Submission
 Submitted (16-MAR-2000) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On May 25, 2000 this sequence version replaced gi:7342114.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/MIT Center for Genome Research
 Center code: WIBR
 Web site: <http://www-seq.wi.mit.edu>
 Contact: sequence_submissions@genome.wi.mit.edu
 Project Information
 Center project name: U7454
 Center clone name: 353_K_12

Summary Statistics
 Sequencing vector: M13: M77815; 100% of reads
 Chemistry: Dye-terminator Big Dye, 100% of reads
 Assembly program: Phrap; version 0.960731

Consensus quality: 156061 bases at least Q40
 Consensus quality: 163284 bases at least Q30
 Consensus quality: 166861 bases at least Q20
 Insert size: 182000; agarose-1P
 Insert size: 169762; sum-of-contigs
 Quality coverage: 4.0 in Q20 bases; agarose-fP
 Quality coverage: 4.3 in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently
 consists of 25 contigs. The true order of the pieces
 is not known and their order in this sequence record is
 arbitrary. Gaps between the contigs are represented as
 runs of N, but the exact sizes of the gaps are unknown.
 This record will be updated with the finished sequence
 as soon as it is available and the accession number will
 be preserved.

1 1105 1204: contig of 1104 bp in length
 * 1105 1204: gap of 100 bp
 * 1205 3226: contig of 2022 bp in length
 * 3227 3326: gap of 100 bp
 * 3327 4615: contig of 1389 bp in length
 * 4616 4715: gap of 100 bp
 * 4716 6570: contig of 1855 bp in length
 * 6571 6670: gap of 100 bp
 * 6671 8592: contig of 1322 bp in length
 * 8593 8692: gap of 100 bp
 * 8693 11929: contig of 3237 bp in length
 * 11930 12029: gap of 100 bp
 * 12030 15549: contig of 3320 bp in length
 * 15550 15549: gap of 100 bp
 * 15650 19338: contig of 3689 bp in length
 * 19339 19438: gap of 100 bp
 * 19439 23177: contig of 3739 bp in length
 * 23178 23277: gap of 100 bp
 * 23278 29032: contig of 5755 bp in length
 * 29033 29132: gap of 100 bp
 * 40698 47278: contig of 6581 bp in length
 * 47279 47778: gap of 100 bp
 * 54797 54796: contig of 7418 bp in length
 * 54897 54996: gap of 100 bp
 * 54897 59726: contig of 4830 bp in length
 * 59727 59826: gap of 100 bp
 * 59827 68926: contig of 9100 bp in length
 * 68927 69026: gap of 100 bp
 * 69027 75941: contig of 6921 bp in length
 * 75948 76047: gap of 100 bp
 * 76048 83423: contig of 7316 bp in length
 * 83424 83523: gap of 100 bp

* 83524 90514: contig of 6991 bp in length
 * 90515 90614: gap of 100 bp
 * 90515 99044: contig of 8420 bp in length
 * 99035 99134: gap of 100 bp
 * 99035 109649: contig of 10515 bp in length
 * 10950 109749: gap of 100 bp
 * 10950 122239: contig of 12550 bp in length
 * 122300 122399: gap of 100 bp
 * 122300 137339: contig of 14970 bp in length
 * 13370 137469: gap of 100 bp
 * 137470 154139: contig of 16730 bp in length
 * 154300 154299: gap of 100 bp
 * 154300 172162: contig of 17863 bp in length

FEATURES
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 Location/Qualifiers
 1..172162
 /organism="Homo sapiens"
 /db_xref="taxon:9666"
 /chromosome="8"
 /map="8"
 /clone_id="RPCI-11_Human_Male_BAC"
 misc_feature
 1..1104
 /note="assembly_fragment"
 misc_feature
 1205..3226
 /note="assembly_fragment"
 misc_feature
 3327..4615
 /note="assembly_fragment"
 misc_feature
 4716..670
 /note="assembly_fragment"
 misc_feature
 6671..8592
 /note="assembly_fragment"
 misc_feature
 8693..11929
 /note="assembly_fragment"
 misc_feature
 12030..15549
 /note="assembly_fragment"
 misc_feature
 15650..19338
 /note="assembly_fragment"
 misc_feature
 19439..23177
 /note="assembly_fragment"
 clone_end:T7
 vector_side:right"
 23278..29032
 /note="assembly_fragment"
 misc_feature
 29133..34839
 /note="assembly_fragment"
 misc_feature
 34897..40597
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 misc_feature
 40698..47778
 /note="assembly_fragment"
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 47379..54796
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 69027..75947
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 misc_feature
 76048..83423
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 misc_feature
 83524..90514
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 90615..90634
 /note="assembly_fragment"
 misc_feature
 99135..109649
 /note="assembly_fragment"
 misc_feature
 109750..122239
 /note="assembly_fragment"
 misc_feature
 122400..137369
 /note="assembly_fragment"
 misc_feature
 137470..154199
 /note="assembly_fragment"

* 11415 11514: gap of 100 bp
 * 14617: contig of 3103 bp in length
 14618 14717: gap of 100 bp
 * 14718 18426: contig of 3709 bp in length
 * 18427 18526: gap of 100 bp
 * 18527 22011: contig of 3495 bp in length
 * 22012 22111: gap of 100 bp
 * 26911: contig of 4807 bp in length
 26919 27018: gap of 100 bp
 * 27019 32120: contig of 5102 bp in length
 * 32121 32220: gap of 100 bp
 * 32221 37187: contig of 4967 bp in length
 * 37188 37287: gap of 100 bp
 * 37288 41477: contig of 4190 bp in length
 * 41478 41577: gap of 100 bp
 * 41578 46736: contig of 5159 bp in length
 * 46737 46836: gap of 100 bp
 * 46837 54164: contig of 7338 bp in length
 * 54165 54244: gap of 100 bp
 * 54265 58419: contig of 4155 bp in length
 * 58420 58519: gap of 100 bp
 * 58520 67538: contig of 9019 bp in length
 * 67539 67638: gap of 100 bp
 * 67639 74093: contig of 6455 bp in length
 * 74094 74193: gap of 100 bp
 * 74194 89734: contig of 1551 bp in length
 * 89735 89834: gap of 100 bp
 * 89835 109850: contig of 20016 bp in length
 * 109851 109950: gap of 100 bp
 * 109951 131927: contig of 21977 bp in length
 * 131928 132027: gap of 100 bp
 * 132028 163683: contig of 31056 bp in length.
 Location/Qualifiers
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 /Organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="1"
 /map="1"
 /clone="RP11-306I4"
 /clone_lib="RPCI-11 Human Male BAC"
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 1..1333
 /note="assembly_fragment"
 misc_feature
 1434..2643
 /note="assembly_fragment"
 misc_feature
 2744..4634
 /note="assembly_fragment"
 misc_feature
 4735..7179
 /note="assembly_fragment"
 misc_feature
 7280..9066
 /note="assembly_fragment"
 misc_feature
 9167..11414
 /note="assembly_fragment"
 misc_feature
 11515..14617
 /note="assembly_fragment"
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 14718..18426
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 18527..22011
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 misc_feature
 22112..26918
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 27019..32120
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 misc_feature
 32221..37187
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 misc_feature
 37288..41477
 /note="assembly_fragment"
 misc_feature
 41578..46736
 /note="assembly_fragment"
 misc_feature
 46837..54164
 /note="assembly_fragment"
 misc_feature
 54265..58419
 /note="assembly_fragment"

AUTHORS	Waterson, R. H.
TITLE	Direct Submission
JOURNAL	Submitted (01-MAR-2000) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis MO 63108, USA
COMMENT	On Jun 17, 2000 this sequence version replaced gi:7263917.
Center: Washington University Genome Sequencing Center	Genome Center
Center code: WUGSC	
Web site: http://genome.wustl.edu/gsc/index.shtml	
Project Information	
Center project name: H_NH0810106	
Summary Statistics	
Sequencing vector: M13; 100%	
Sequencing vector: Plasmid; 0%	
Chemistry: Dye-primer ET; 97% of reads	
Chemistry: Dye-terminator Big Dye; 3% of reads	
Assembly program: Phrap; version 0.990319	
Consensus quality: 173830 bases at least Q40	
Consensus quality: 175557 bases at least Q30	
Consensus quality: 176446 bases at least Q20	
Insert size: 242000; agarose-fp	
Insert size: 177938; sum-of-contigs	
Quality coverage: 5.32 in Q20 bases; agarose-fp	
Quality coverage: 6.01 in Q20 bases; sum-of-contigs	
* NOTE: This is a 'working draft' sequence. It currently consists of 17 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.	
* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.	
* 1 1801: contig of 1801 bp in length	
* * 1901: gap of unknown length	
* * 1902 4641: contig of 2740 bp in length	
* * 4642 4741: gap of unknown length	
* * 4742 9679: contig of 4938 bp in length	
* * 9680 9779: gap of unknown length	
* * 9780 13392: contig of 3613 bp in length	
* * 13393 13492: gap of unknown length	
* * 13493 17164: contig of 3672 bp in length	
* * 17165 17264: gap of unknown length	
* * 17265 21510: contig of 4246 bp in length	
* * 21511 21610: gap of unknown length	
* * 21611 27089: contig of 5479 bp in length	
* * 27090 27189: gap of unknown length	
* * 27190 35636: contig of 8447 bp in length	
* * 35637 35736: gap of unknown length	
* * 35737 45537: contig of 9801 bp in length	
* * 45538 45637: gap of unknown length	
* * 45638 56451: contig of 10814 bp in length	
* * 56452 56551: gap of unknown length	
* * 56552 66802: contig of 10251 bp in length	
* * 66803 66902: gap of unknown length	
* * 66903 79118: contig of 12216 bp in length	
* * 79119 79218: gap of unknown length	
* * 79219 94355: contig of 15137 bp in length	
* * 94356 94455: gap of unknown length	
* * 94456 111274: contig of 16819 bp in length	
* * 111275 111374: gap of unknown length	
* * 111375 130600: contig of 19226 bp in length	
* * 130601 130700: gap of unknown length	
* * 130701 150278: contig of 19578 bp in length	
* * 150279 150378: gap of unknown length	
* * 150379 179538: contig of 29160 bp in length.	
Location/Qualifiers	
1. 119538 /organism="Homo sapiens"	
/db_xref="Taxon:9606"	
/chromosome="19"	
FEATURES	
source	

Qy	7273	tgactgtggccacaatctgttgttcaggcagatccatccatcgct	7324
LOCUS	AL355392	159440 bp	DNA
DEFINITION	Human DNA sequence from clone RP5-1187J4 on chromosome 20q11.1-11.21, complete sequence.	13-SEP-2000	PRI
VERSION	AL355392.7	GI:10178502	
KEYWORDS	HGVS		
SOURCE	Homo sapiens		
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1 (bases 1 to 159440)		
AUTHORS	Bird,C.		
TITLE	Direct Submission		
JOURNAL	Submitted (12-SEP-2000) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk		
REQUESTS	clonerequest@sanger.ac.uk		
COMMENT	On Sep 15, 2000 this sequence version replaced Gi:9213549. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em', EMBL; Sw', SWISSPROT; Tr', TREMBL; Wp', WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 20, constructed by the Sanger Centre Chromosome 20 Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/Chr20 RP5-1187J4 is from the library RCI-5 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see http://bacpac.med.buffalo.edu/VECTOR: PCPAC2 This sequence is the entire insert of clone RP5-1187J4. The true left end of clone RP1-327D19 is at 62286 in this sequence.		
FEATURES	Location/Qualifiers		
source	1..159440		
	/organism="Homo sapiens"		
	/db_xref="taxon:9606"		
	/chromosome="20"		
	/map="q11.1-11.23"		
	/clone_id="RP5-1187J4"		
	/clone_idb="RCI-5"		
repeat_region	816..1129		
	/note="AlusX repeat: matches 1..312 of consensus"		
	/note="AlusX repeat: matches 1..311 of consensus"		
repeat_region	1207..1334		
	/note="AlusX repeat: matches 64..175 of consensus"		
repeat_region	1335..1848		
	/note="137 copies 2 mer ga 61% conserved"		
misc_feature	1757..1924		
	/note="Random repeat. weak data"		
repeat_region	1917..2031		
	/note="MRB repeat: matches 48..160 of consensus"		
repeat_region	2415..2622		
	/note="MRB repeat: matches 20..253 of consensus"		
repeat_region	2810..3103		
	/note="AlusX repeat: matches 1..296 of consensus"		


```

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OM nucleic - nucleic search, using sw model
Run on: April 5, 2001, 19:21:50 ; Search time 4648.59 Seconds
          (without alignments)
          13639.351 Million cell updates/sec

Title: US-09-513-888-1
Perfect score: 9048
Sequence: 1 gccttccaaagacctgccc.....tgccatttcacgccccct 9048

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 7991742 seqs, 3503743858 residues

Total number of hits satisfying chosen parameters:
Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
                  Maximum Match 100%
Listing first 45 summaries

Database : EST:*
1: qb_est1:*
2: qb_est2:*
3: qb_est3:*
4: qb_est4:*
5: qb_est5:*
6: qb_est6:*
7: qb_est7:*
8: qb_est8:*
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11: qb_est11:*
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45: em_esthum3:*
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103: qb.est57:*
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112: em.esthum21:*
113: em.esthum22:*
114: em.esthum23:*
115: em.estom1:*
116: em.estom2:*

```

```

117: em_estpl6;*
118: em_estpl7;*
119: em_estpl8;*
120: em_estpl9;*
121: em_estrl5;*
122: em_estrl6;*
123: em_estrl7;*
124: em_estrl8;*
125: em_estrl9;*
126: gb_est38;*
127: gb_est59;*
128: gb_est60;*
129: gb_est61;*
130: gb_est62;*
131: gb_est3;*
132: gb_est64;*
133: gb_est65;*
134: gb_est66;*
135: gb_est75;*
136: gb_est76;*
137: gb_est77;*
138: gb_est78;*
139: gb_est79;*
140: gb_est80;*
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142: gb_est82;*
143: gb_est33;*
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145: gb_est85;*
146: gb_est86;*
147: gb_est87;*
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149: gb_gss2;*
150: gb_gss3;*
151: gb_gss4;*
152: em_gss1;*
153: em_gss2;*
154: em_gss3;*
155: em_gss4;*
156: gb_gss5;*
157: gb_gss6;*
158: gb_gss7;*
159: gb_gss8;*
160: gb_gss9;*
161: gb_gss10;*
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163: gb_gss12;*
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177: em_gss6;*
178: em_gss8;*
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180: em_gss10;*
181: em_gss11;*
182: em_gss12;*
183: em_gss13;*
184: em_gss14;*
185: em_gss15;*
186: em_gss16;*
187: em_gss17;*
188: em_gss18;*
189: em_gss19;*

```

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB	ID	Description
1	704.2	7.8	705	28	AJ118597	DKEZP761D	N31948 YY22912.s1
c	2	549.8	6.1	599	142	N31948	AW137540 UI-H-BI1-
c	3	525.2	5.8	547	40	AW007737	wt6806.x
c	4	524.4	5.8	545	38	AW590735	h948h03.x
c	5	521.2	5.8	529	92	AW590735	N21184 YX41a10.s1
c	6	502.4	5.6	619	142	N21184	YX41a10.s1
c	7	484.4	5.4	817	27	AJ984777	wR8507.x
c	8	480.8	5.3	743	107	BE10921	601303579
c	9	468.8	5.2	472	28	AL135755	DKEZP762B
c	10	468.4	5.2	487	19	AI357233	Q6304.0
c	11	468	5.2	673	107	BE384131	601272956
c	12	447.2	4.9	452	39	AW051159	N35845 YX89f10.r1
c	13	445.4	4.9	447	15	AI042490	0x6204.x
c	14	440.4	4.9	442	19	AI360882	QY01c12.x
c	15	439.8	4.9	453	23	AI633596	th70a03.x
c	16	439.4	4.9	451	23	AI636674	ts92e04.x
c	17	433.6	4.8	643	142	N35845	YX89f10.r1
c	18	427.8	4.7	432	38	AW002410	wu6105.x
c	19	423.4	4.7	425	14	BE049448	t800408.x
c	20	421.4	4.7	434	22	AI62326	ts800408.x
c	21	416.4	4.6	418	15	AI078630	0x51b09.x
c	22	403.6	4.5	495	142	NA2784	YY22912.r1
c	23	401.4	4.4	796	106	BE279166	BE279166
c	24	392.4	4.3	450	23	AI652496	wb29102.x
c	25	391.8	4.3	401	19	AI362152	Q441b07.x
c	26	387	4.3	558	107	BE34676	BE34676
c	27	385	4.3	461	142	NA2784	YY22912.r1
c	28	368.8	4.1	531	106	BE276168	BE276168
c	29	365	4.0	397	39	AW028197	AW028197
c	30	362.4	4.0	797	109	BE510725	BE510725
c	31	351.4	3.9	463	1	AA020852	z64908.r
c	32	342.6	3.8	471	141	H09157	H09157
c	33	342.6	3.8	751	106	BE312985	BE312985
c	34	336.6	3.7	344	38	AW016544	UI-H-BI0P
c	35	335	3.7	357	15	AI080440	0x8c12.s
c	36	303	3.3	328	23	AI636335	tzz78a05.x
c	37	300	3.3	323	23	AI636335	F1182 HSC32a091.n
c	38	299.4	3.3	302	140	T65388 YC73b091.n	R88008 Ym87502.r1
c	39	299.2	3.3	349	145	T65388	Ym87502.r1
c	40	299.2	3.3	448	144	R88008	Ym87502.r1
c	41	296.2	3.3	326	28	AL047147	AL047147
c	42	296	3.3	296	28	AL134288	DKEZP547J
c	43	279	3.1	347	144	R87502	R87502.s1
c	44	257.4	2.8	274	140	F09471	F09471 HSC32a092.n
c	45	254.8	2.8	407	38	AV663617	AV663617

ALIGNMENTS

RESULT	1	LOCUS	ALL18597	705 bp	mRNA
DEFINITION		DKEZP761D0110-r1	761	(synonym: hamy2)	EST
ACCESSION		DKEZP761D0110-5;		mRNA sequence	Homo sapiens cDNA clone
VERSION		ALL18597			
KEYWORDS		EST			
SOURCE		human			

Page 6

from the same library (cloneIDs 1257096-1258631, 146906-1470983, and 147552-147643). Subtraction by Bento Soares and M. Fatima Bonaldo. " 141 a 146 c 134 g 108 t

High quality sequence stops : 342
Source: IMAGE Consortium, LNL
This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@lnl.gov) for further information.
Insert length: 2372 std. error: 0.00

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Copyright (c) 1993 - 2000 Compugen Ltd.		GenCore version 4.5	
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Perfect score:	9048		
Sequence:	1 gccttccaaggacctgtcccc.....tgccatttctcacggccctct 9048		
Scoring table:	IDENTITY NUC		
	Gapop 10.0 , Gapext 1.0		
Searched:	480022 seqs, 187831343 residues		
Total number of hits satisfying chosen parameters:	960044		
Post-processing: Minimum Match 10%			
	Maximum Match 10%		
	Listing first 45 summaries		
Minimum DB seq length:	0		
Maximum DB seq length:	20000000000		
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	9: /cgcn2_2/gcdatata/geneseq/geneseq/geneseq/NNA1988.DAT:*		
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	19: /cgcn2_2/gcdatata/geneseq/geneseq/geneseq/NNA1998.DAT:*		
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Pred. No.	is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.		
SUMMARIES			
Result No.	Score	Query Match Length	DB ID Description
1	365.8	4.0	393 16 T23583 Human gene signature
c 2	212.2	2.3	700 20 X30339 DNA encoding a hum
c 3	211.2	2.3	54548 21 245596 DNA sequence of th
c 4	209.9	2.3	2116 20 X87623 Novel CD24* ORF en
c 5	209.9	2.3	2194 20 277555 Human ovarian tumo
c 6	209.8	2.3	2233 20 224431 Human bladder tumo
c 7	206.8	2.3	45556 20 X23520 Human kidney amino
c 8	205.8	2.3	9365 21 250359 Human CD39-L4 geno
9	205.4	2.3	138169 21 A34791 Human adenosine re
10	205.4	2.3	141589 21 A35005 Human adenosine re
11	205.4	2.3	141589 21 A35030 Human adenosine re
c 12	204.2	2.3	1601 21 A35191 Human adenosine re
SUMMARIES			
Result No.	Score	Query Match Length	DB ID Description
1	365.8	4.0	393 16 T23583 Human gene signature
c 2	212.2	2.3	700 20 X30339 DNA encoding a hum
c 3	211.2	2.3	54548 21 245596 DNA sequence of th
c 4	209.9	2.3	2116 20 X87623 Novel CD24* ORF en
c 5	209.9	2.3	2194 20 277555 Human ovarian tumo
c 6	209.8	2.3	2233 20 224431 Human bladder tumo
c 7	206.8	2.3	45556 20 X23520 Human kidney amino
c 8	205.8	2.3	9365 21 250359 Human CD39-L4 geno
9	205.4	2.3	138169 21 A34791 Human adenosine re
10	205.4	2.3	141589 21 A35005 Human adenosine re
11	205.4	2.3	141589 21 A35030 Human adenosine re
c 12	204.2	2.3	1601 21 A35191 Human adenosine re

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

ALIGMENTS

RESULT 1

T23583 ID T23583 standard; CDNA to mRNA; 393 BP.

XX AC T23583;

XX DT 02-SEP-1996 (first entry)

XX DE Human gene signature HUMGS05436.

XX KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;

XX KW human; cloning; mapping; non-biased library; diagnosis; detection;

XX KW cell typing; abnormal cell function; ss.

XX OS Homo sapiens.

XX PN WO9514772-A1.

XX PD 01-JUN-1995.

XX PF 11-NOV-1994; 94WO-JP01916.

XX PR 12-NOV-1993; 93JP-0355504.

PA (MATS/) MATSUBARA K.

PA (OKUB/) OKUBO K.

XX PI Matsubara K, Okubo K;

XX DR WPI; 1995-206931/27.

XX PT Identifying gene signatures in 3'-directed human cDNA library - e.g. PT for diagnosis of abnormal cell function, by preparing cDNA that PT reflects relative abundance of corresp. mRNA in specific human tissues


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Db	1343	TGAAACCTCTGTCTACTAATTAATACAAAAATTAGCCGGCANGTTGCAGGTGCTGTA	1284
Qy	4479	atccccagtcacttggggggctggcaggagaatc-----aggggggggggggggttg	4532
Db	1283	AATCCAGCTACTGTGGGGTGAAGAGAAATGGCTGAGCCGGGCAAATGGTA	1224
Qy	4533	cagttagccaaggatcaaggccactacaccggccatggttgacaaggcggactttccaa	4592
Db	1223	CACTGACTGAGATCGACCACCTGACTCGGGCTGGCGACAAAGTGAGACITGCTAA	1164
Qy	4593	tattaaaccaaataataactatgtgtcattatacatgtgattttatccat	4651
Qy	1163	AAATAATAATAAAAAGCATTTGAAATTAGTCGGTCAATGCCAATCTACTCTT	1105
RESULT 7			
X	x3520/c	X3520 standard; DNA: 45546 BP.	
XX	XX	XX 23-JUN-1999 (first entry)	
DE	DE	Human kidney aminopeptidase P genomic DNA fragment 4.	
XX	XX	XX Aminopeptidase; human; Amp; gene therapy; treatment; Amp-deficiency; prenatal diagnosis; angioedema; antihypertensive agent; atherosclerosis; arterial stenosis; industrial protein feed; malabsorption syndrome; proteinaceous waste degradation; additive; immunohistochemistry; ss.	
XX	XX	XX Homo sapiens	
OS	OS	XX WO911799-A2.	
XX	XX	XX PD 11-MAR-1999.	
XX	XX	XX PF 02-SEP-1998; 98WO-US18426.	
XX	XX	XX PR 02-SEP-1997; 97US-0057854.	
XX	XX	XX (MEDI-) MEDICAL COLLEGE GEORGIA RES INST.	
PA	PA	XX Ryan JW, Sprinkle TJC, Venema RC;	
PI	PI	XX DR 1999-205193/17.	
XX	XX	XX Nucleic acid encoding human aminopeptidase P	
PT	PT	XX Claim 13; Page 165-192; 201pp; English.	
PS	PS	XX CC This invention describes the isolation of a novel human aminopeptidase (AMP). This protein is used to produce recombinant AMP and can be used for gene therapy for treating Amp-deficiency conditions. Its fragments are used as primers and probes to identify patients with homozygous and heterozygous AMP deficiency, including prenatal diagnosis (patients affected in AMP are at risk of developing angioedema, also as antisense inhibitors of angiotensin converting enzyme inhibitors). The product of the invention is also used to identify Amp-expressing sequences in other animals and to generate transgenic animals, and comparisons of genomic sequences are used to detect mutations. AMP inhibitors are potentially useful as antihypertensive agents and to prevent or treat arterial (re)stenosis or atherosclerosis. The structure of AMP is used to design synthetic substrates, e.g. for use in AMP assays. AMP, which hydrolyzes N-terminal imido bonds, can be used to degrade industrial protein feeds to free amino acids, to degrade proteinaceous wastes, as additives in enzyme formulations used to treat malabsorption syndrome and for studying its biological role. Antibodies against AMP are used in immunohistochemical methods to study AMP distribution.	

nucleic acids involved in bronchoconstriction, allergies, and/or inflammation. The ON can have antiinflammatory, antiallergic, antiasthmatics, cyrostatic and analgesic activities. The compositions are useful for the treatment of diseases associated with inflammation, impaired airways, including lung disease and diseases whose secondary effects afflict the lungs of a subject. They can be used for treating e.g. ischaemic conditions, pulmonary vasoconstriction, allergies, asthma, impeded respiration, respiratory distress syndrome, pain, cystic fibrosis, pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), and cancers such as leukaemias, lymphomas, carcinomas, and cancers which may metastasise to the lungs, including breast and prostate cancer. The reduction of the adenosine content of the ONs reduces side effects. The A-containing ONs break down with the release of deoxyadenosine which activates adenosine receptors causing bronchoconstriction and inflammation. A32313 to A35312 represent the nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 2815, and then the last 185 sequences are also called SEQ ID NO:1 to 185, but the sequences differ from the previously named sequences. SEQ ID NO:11 to 1680 (A32323 to A33992) are specifically claimed ONs from the present invention. N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequences given in the sequence listing.

Sequence 141589 BP; 42856 A; 28938 C; 26863 G; 42932 T; 0 other;

cancer; leukaemia; lymphoma; carcinoma; metastasis; ss
Homo sapiens.
WO200009525-A2.
24-FEB-2000. 99WO-US17712.
03-AUG-1999; 98US-0095212.
03-AUG-1998;
(UYEC-) UNIV EAST CAROLINA.
Nyce JW;
WPI: 2000-205971/18.
New antisense oligonucleotides useful for treating e.g. vasocstriction, inflammation, allergies, asthma, hypertension, bronchitis, emphysema, respiratory distress syndrome, i - cancers -

oligonucleotide (ON) with low adenosine (up to 15%), which targets nucleic acids involved in bronchoconstriction, allergies, and/or inflammation. The ON can have antiinflammatory, antiallergic, antiasthmatic, cytostatic and analgesic activities. The compositions are useful for the treatment of diseases associated with inflammation, impaired airways, including lung disease and diseases whose secondary effects afflict the lungs of a subject. They can be used for treating e.g. ischaemic conditions, pulmonary vasodilator syndrome, asthma, impaired respiration, respiratory distress syndrome, pain, cystic fibrosis, pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), and cancers such as leukaemias, lymphomas, carcinomas, and cancers which may metastasise to the lungs, including breast and prostate cancer. The reduction of the adenosine content of the ONs reduces side effects. The A-containing ONs break down with the release of deoxyadenosine which activates adenosine receptors causing bronchoconstriction and inflammation. A32313 to A35312 represent the nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 2815, and then the last 185 sequences are also called SEQ ID NO:1 to 185, but the sequences differ from the previously named sequences. SEQ ID NO:1 to 1680 (A33323 to A33992) are specifically claimed ONs from the present invention.

N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequences given in the sequence listing.

SQ Sequence 1601 BP; 445 A; 368 C; 356 G; 432 T; 0 other

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GenCore - nucleic search, using sw model

on on: April 5, 2001, 19:35:45 ; Search time 152.82 Seconds
(without alignments)

9541.804 Million cell updates/sec

title: US-09-5113-888-1

perfect score: 9048

Sequence: 1 gctttccaaagccctgtcccc.....tgccatttctcacggccct 9048

scoring table: IDENTITY_NUC Gapo 10.0 , Gapext 1.0

searched: 280836 seqs, 80580151 residues

Total number of hits satisfying chosen parameters: 561672

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 100%
Maximum Match 100%
Listing first 45 summaries

Database : Issued_Patents_NA.*

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5: /egn2_6/podata/2/ina/backfile1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

result No.	Score	Query Match	Length	DB ID	Description
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C 2	209.8	2.3	35060	3 US-08-814-095-7	Sequence 7, Appli
C 3	201.6	2.2	53526	3 US-08-658-136-2	Sequence 2, Appli
C 4	201.6	2.2	53577	3 US-08-658-136-1	Sequence 1, Appli
C 5	199	2.2	1988	4 PCT-US95-07201-11	Sequence 11, Appli
C 6	199	2.2	22481	4 PCT-US95-07201-43	Sequence 43, Appli
C 7	199	2.2	1600	2 US-08-487-088-1	Sequence 117, App
C 8	195.8	2.2	1600	2 US-08-487-088-1	Sequence 117, App
C 9	195.8	2.2	1600	2 US-08-487-088-1	Sequence 117, App
C 10	195.8	2.2	7210	2 US-08-25-963B-10	Sequence 10, Appli
C 11	195.8	2.2	7210	4 PCT-US95-07201-10	Sequence 10, Appli
C 12	195.8	2.2	22481	4 PCT-US95-07201-43	Sequence 43, Appli
C 13	195.2	2.2	7676	1 US-08-45-777A-7	Sequence 7, Appli
C 14	195.2	2.2	7676	2 US-08-45-777A-7	Sequence 7, Appli
C 15	195.2	2.2	7676	2 US-08-720-420A-117	Sequence 7, Appli
C 16	195.2	2.2	7676	4 PCT-US95-06743-7	Sequence 7, Appli
C 17	194	2.1	20303	1 US-08-370-975B-6	Sequence 6, Appli
C 18	194	2.1	26764	1 US-08-370-975B-1	Sequence 1, Appli
C 19	194	2.1	87350	3 US-08-781-891-79	Sequence 79, Appli
C 20	193.8	2.1	282	1 US-08-133-629-8	Sequence 8, Appli
C 21	193.6	2.1	4823	2 US-08-45-254-5	Sequence 5, Appli
C 22	193.6	2.1	4823	2 US-08-48-257-20	Sequence 20, Appli
C 23	193.6	2.1	4823	3 US-08-993-927-5	Sequence 5, Appli
C 24	193.6	2.1	4823	4 PCT-US94-08806-28	Sequence 28, Appli
C 25	193.6	2.1	4823	4 PCT-US95-01829-5	Sequence 5, Appli
C 26	193.6	2.1	4823	4 PCT-US95-16626-5	Sequence 5, Appli
C 27	193.2	2.1	4220	2 US-08-93-983-66	Sequence 66, Appli
C 28	193.2	2.1	4220	2 US-08-83-877-66	Sequence 66, Appli

ALIGNMENTS

RESULT 1
US-08-848-252-1/c

Sequence 1, Application US/08848232

Patent No. 5804177

GENERAL INFORMATION:

APPLICANT: Humphries, Keith R.

TITLE OF INVENTION: METHOD OF USING CD24 AS A CELL MARKER

NUMBER OF SEQUENCES: 4

CORRESPONDENCE ADDRESS:

ADDRESSEE: Bereskin & Parr

STREET: 40 King Street West

CITY: Toronto

STATE: Ontario

COUNTRY: Canada

ZIP: M5H 2Y2

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/848-252

FILING DATE: 29-APR-1997

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/538,052

FILING DATE:

APPLICATION NUMBER: US 08/151,672

ATTORNEY/AGENT INFORMATION:

NAME: McDairmid, Shona S.

REGISTRATION NUMBER: P-38,798

REFERENCE/DOCKET NUMBER: 3158-028

TELECOMMUNICATION INFORMATION:

TELEPHONE: (416) 364-7311

TELEFAX: (416) 361-1398

TELEX: 06-23115

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1811 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

ORIGINAL SOURCE:

ORGANISM: Homo sapiens

IMMEDIATE SOURCE:

LIBRARY:

CLONE: Signal Transductor CD24

NAME/KEY: sig-peptide

Page 3

Qy 7178 cagatttttttt 7189
 Db 840 CAGAAATACTT 851

RESULT 7
PCT-US95-07201-43

; GENERAL INFORMATION:
 / APPLICANT: Chader, Gerald J.; Becerra, Sofia
 / APPLICANT: Patricia; Schwartz, Joan P.;
 / APPLICANT: Tanikaki, Takayuki
 / TITLE OF INVENTION: PIGMENT EPITHELIUM
 / TITLE OF INVENTION: DERIVED FACTOR: CHARACTERIZATION GENOMIC
 / TITLE OF INVENTION: ORGANIZATION AND SEQUENCE OF THE PEDF GENE
 / NUMBER OF SEQUENCES: 43
 / CORRESPONDENCE ADDRESS:
 / ADDRESSEE: Morgan & Finnegar, L.L.P.
 / STREET: 345 Park Avenue
 / STATE: New York
 / COUNTRY: USA
 ZIP: 10154

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy Disk
 COMPUTER: IBM PC Compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: WORDPERFECT 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: PCT/US95/07201
 FILING DATE: 06-JUN-1995
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/367,841
 FILING DATE: 30-DEC-1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/257,963
 FILING DATE: 07-JUN-1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 07/952,796
 FILING DATE: 22-SEP-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: DOROTHY R. AUTH
 REGISTRATION NUMBER: 36434
 REFERENCE/DOCKET NUMBER: 20264126PCT

TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212) 758-4800
 TELEFAX: (212) 751-6849

INFORMATION FOR SEQ ID NO: 43:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 22481 Base Pairs
 TYPE: Nucleic Acid
 STRANDEDNESS: Double
 TOPOLOGY: Unknown
 MOLECULE TYPE: Genomic DNA
 FEATURE:
 NAME/KEY: P1-147
 LOCATION:
 IDENTIFICATION METHOD:
 OTHER INFORMATION: full length genomic
 PCT-US95-07201-43

Qy Match 2.2%; Score 199; DB 4; Length 22481;
 Best Local Similarity 78.8%; Pred. No. 6e-33;
 Matches 246; Conservative 0; Mismatches 65; Indels 1; Gaps 1;

Db 6878 tttattttaattttatgagacagggtctgtcccccaggctggatcgatgg 6937
 Db 15138 TNTTTTNNNNCTTCGACGGAGTCCTTGNCAGGCTGAGTCAGTGG 15197
 Qy ~6938 catgatcatagtcactgcagccatactcctggctcaagcaaatccctgcctcagc 6937

RESULT 8

US-08-487-113D-117/C

; Sequence 117, Application US/08487113D
 ; Patent No. 5837822
 / GENERAL INFORMATION:
 / APPLICANT: Galatin, W. Michael
 / APPLICANT: Vaeux, Rosemary
 / TITLE OF INVENTION: ICAM-Related Materials and Methods
 / NUMBER OF SEQUENCES: 120
 / CORRESPONDENCE ADDRESS:
 / ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
 / STREET: 6300 Sears Tower, 233 South Wacker Drive
 / CITY: Chicago
 / STATE: Illinois
 / COUNTRY: United States of America
 / ZIP: 60606-5402
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC DOS/MS-DOS
 SOFTWARE: Patent Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/487,113D
 / FILING DATE:
 / CLASSIFICATION: 424
 / PRIOR APPLICATION DATA:
 / APPLICATION NUMBER: US 08/286,754
 / FILING DATE: 05-AUG-1994
 / PRIOR APPLICATION DATA:
 / APPLICATION NUMBER: US 08/102,852
 / FILING DATE: 05-AUG-1993
 / PRIOR APPLICATION DATA:
 / APPLICATION NUMBER: US 08/4894,061
 / FILING DATE: 05-JUN-1992
 / PRIOR APPLICATION DATA:
 / APPLICATION NUMBER: US 08/009,266
 / FILING DATE: 22-JAN-1993
 / PRIOR APPLICATION DATA:
 / APPLICATION NUMBER: US 07/894,061
 / FILING DATE: 27-JAN-1992
 / ATTORNEY/AGENT INFORMATION:
 / NAME: No. 5637822 and, Greta E.
 / REGISTRATION NUMBER: 35,302
 / REFERENCE/DOCKET NUMBER: 32744
 / TELEPHONE: (312) 474-6300
 / TELEX: 25-356
 / INFORMATION FOR SEQ ID NO: 117:
 / SEQUENCE CHARACTERISTICS:


```

; TELECOMMUNICATION INFORMATION:
;   TELEPHONE: 610-270-5364
;   TELEFAX: 610-270-5090
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7676 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US - 08-998-208 7

Query Match 2.2%; Score 195.2; DB 2; Length 7676;
Best Local Similarity 80.5%; Pred. No. 2.4e-32;
Matches 243; Conservative 0; Mismatches 53; Indels 6; Gaps 1;
Qy 4312 tggtggcttaggcgtggctcacgccctgcaatcccaaggacttggggggcaggccg 4371
Db 6063 TGAATGGTGGCCGGGCTACACCTGTAAATCCAGCACTTGGAAAGCTGGCAG 6004
Qy 4372 gceagggacttggctggatcgatcatcgatccatgtggaaacctcccgct 4431
Db 6003 GCACATTGCCTGAGTCAGGAGCTGAGGCCAACATGGGAAACCCCGCT 5944
Qy 4432 ctactaaaaataaaaaatccaggcaggcatggttggcaatgtctgttaatccccactt 4491
Db 5943 CTACTAAAAATAACAAAAGTTAGCCCCGATCCGGCATCCGGCTGTATCCAGCTACTC 5884
Qy 4492 ggaggctggggggat-----cagaggggggggggggatgtggatggccaga 4545
Db 5883 AGGGGGCTGGGAGGAGATGCTGAGCTGGGGGAGTTGCACTGGAGTTGAGTGGAGA 5824
Qy 4546 tcacggccactacccggccatggctggccaaatggccatgtggccaaatattacaatataat 4605
Db 5823 TCATGCCACTGCACTTCAGCTGGTACAGTGAGGTGAGGGTCCCTCTCAAAAAAAAG 5764
Qy 4606 aa 4607
Db 5763 AA 5762

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Search completed: April 6, 2001, 00:19:59
 Job time: 17054 sec

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Fri Apr 6 08:40:34 2001

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OM nucleic - nucleic search, using sw model

Run on: April 5, 2001, 19:52:25 ; Search time 318.41 Seconds
(without alignments)

Title: US-09-513-888-1
Perfect score: 9048
Sequence: 1 gccattccaggaccctgcccc.....tgcacatttttcacccctct 9048

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 480022 seqs, 187831343 residues

Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_36 : *
 :

RESULT 1
T23-83 ID T23583 standard; CDNA to mRNA; 393 BP.

XX AC T23583;
XX DT 02-SEP-1996 (first entry)

XX DE Human gene signature HUMGS05436.
XX Gene signature; messenger RNA; mRNA; relative abundance; frequency;
KW human; Cloning; mapping; non-biased library; diagnosis; detection;
KW cell typing; abnormal cell function; ss.
XX OS Homo sapiens.

XX PN WO9514772-A1.
XX PD 01-JUN-1995.

XX PF 11-NOV-1994; 94WO-JP01916.
XX PR 12-NOV-1993; 93JP-0355504.
XX PA (MATS/) MATSUBARA K.
PA (OKIB/) OKUBO K.
XX PI Matsubara K, Okubo K;
XX DR WPI: 1995-206931/27.

XX PT Identifying gene signatures in 3'-directed human cDNA library - e.g.,
PT for diagnosis of abnormal cell function, by preparing cDNA that
PT reflects relative abundance of corresp. mRNA in specific human
tissues

xx Ps Claim 1; Page 1404; 2245pp; Japanese.

xx A single-stranded DNA (or its complementary strand or the corresp. double-stranded DNA) which comprises one of the 7837 "GS" sequences given in WO9001-T26837, and which is able to hybridise to part of human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature) sequences were obtained from 3'-directed cDNA libraries prepared from various human tissues; synthesis of cDNA was initiated from the 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-untranslated sequence is unique to a particular mRNA species, almost all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library is constructed so as to reflect accurately the relative abundance of different mRNAs in the particular tissue from which it was derived. The apparent frequency of given GS in a cDNA library can be determined (esp. using primers and probes derived from the GS sequences) as a means of diagnosing abnormal cell function or for recognising different cell types.

xx Sequence 393 BP; 73 A; 117 C; 121 G; 78 T; 4 other;

Query Match Score 365.8; DB 16; Length 393;
Best Local Similarity 97.2%; Pred. No. 2.1e-56;
Matches 381; Conservative 0; Mismatches 10; Indels 1; Gaps 1;
Qy 8454 gatcggaactctggaccctggaaacctggactgtacgggttccctcttagct 8513
Db 1 gatcgactcgccctggaaacctggactgtacggactgtacgggttacccctcttagct 60
Qy 8514 ctcccaactgtccaggcacacacagccatacgactgtacggactgtacggactgtacgggttacccctcttagct 61
Db 61 ctcccaactgtccaggcacacacagccatacgactgtacggactgtacggactgtacgggttacccctcttagct 120
Qy 8574 ggatatggggcatcttcctccaggaggactcgatgtacccctgtgcctggccc 8633
Db 121 ggataatggggcatcttcctccaggaggactcgatgtacccctgtgcctggccc 180
Qy 8634 agcttggccatcttccttagtgagacatgtggccgaaacttaaggccaggctggctggagg 8693
Db 181 agcttggccatcttccttagtgagacatgtggccgaaacttaaggccaggctggctggagg 240
Qy 8654 agagcgcttggagaaaggattccctcgaggactcatgcgtggggct 8753
Db 241 agacgcgacttggagaaaggattccctcgaggactcatgcgtggggct 300
Qy 8754 gcatgacatggctggacttccttcctggaggacttcacccaccttgtggaaagg 8813
Db 301 gcatgacatggctggacttccttcctggaggacttcacccaccttgtggaaagg 360
Qy 8814 tggccctttgtgc-tgcgcctgtgcaggcaggact 8844
Db 361 tggccctttgtgc-tgcgcctgtgcaggcaggact 392

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